

<b>TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED/ELECTED OFFICE (DO/EO/US) CONCERNING A FILING UNDER 35 U.S.C. 371</b>		<b>Attorney's Docket No.</b> 288748.0005
		<b>U.S. Application No.</b> <b>10/019229</b>
<b>International Application No.</b> PCT/GB00/02513	<b>International Filing Date</b> June 29, 2000	<b>Priority Date Claimed</b> June 29, 1999
<b>Title of Invention:</b> PYRAZOLIDINOL COMPOUNDS		
<b>Applicant(s) For DO/EO/US:</b> Enok TJØTTA; and Jo KLAVENESS		
Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:		
<ol style="list-style-type: none"> <li>1. <input checked="" type="checkbox"/> This is a FIRST submission of items concerning a filing under 35 U.S.C. 371.</li> <li>2. <input type="checkbox"/> This is a SECOND or SUBSEQUENT submission of items concerning a filing under 35 U.S.C. 371.</li> <li>3. <input checked="" type="checkbox"/> This is an express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39(1).</li> <li>4. <input checked="" type="checkbox"/> A proper Demand for International Preliminary Examination was made by the 19<sup>th</sup> month from the earliest claimed priority date.</li> <li>5. <input checked="" type="checkbox"/> A copy of the International Application as filed (35 U.S.C. 371(c)(2))           <ol style="list-style-type: none"> <li>a) <input type="checkbox"/> is transmitted herewith (required only if not transmitted by the International Bureau).</li> <li>b) <input checked="" type="checkbox"/> has been transmitted by the International Bureau. (<b>courtesy copy enclosed</b>)</li> <li>c) <input type="checkbox"/> is not required, as the application was filed in the United States Receiving Office (RO/US).</li> </ol> </li> <li>6. <input type="checkbox"/> A translation of the International Application into English (35 U.S.C. 371(c)(2)).</li> <li>7. <input checked="" type="checkbox"/> A copy of the International Search Report. (PCT/ISA/210).</li> <li>8. <input checked="" type="checkbox"/> Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3))           <ol style="list-style-type: none"> <li>a) <input type="checkbox"/> are transmitted herewith (required only if not transmitted by the International Bureau).</li> <li>b) <input checked="" type="checkbox"/> have been transmitted by the International Bureau. (<b>courtesy copy enclosed</b>)</li> <li>c) <input type="checkbox"/> have not been made; however, the time limit for making such amendments has NOT expired.</li> <li>d) <input type="checkbox"/> have not been made and will not be made.</li> </ol> </li> <li>9. <input type="checkbox"/> A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).</li> <li>10. <input checked="" type="checkbox"/> An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)). (<b>unexecuted</b>)</li> <li>11. <input checked="" type="checkbox"/> A copy of the International Preliminary Examination Report (PCT/IPEA/409).</li> <li>12. <input type="checkbox"/> A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).</li> </ol>		
<b>Items 13. to 20. below concern other document(s) or information included:</b>		
<ol style="list-style-type: none"> <li>13. <input type="checkbox"/> An Information Disclosure Statement under 37 CFR 1.97 and 1.98.</li> <li>14. <input type="checkbox"/> An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.</li> <li>15. <input checked="" type="checkbox"/> A FIRST preliminary amendment.</li> <li>16. <input type="checkbox"/> A SECOND or SUBSEQUENT preliminary amendment.</li> <li>17. <input type="checkbox"/> A substitute specification.</li> <li>18. <input type="checkbox"/> A change of power of attorney and/or address letter.</li> <li>19. <input type="checkbox"/> Certificate of Mailing by Express Mail.</li> <li>20. <input checked="" type="checkbox"/> Other items or information           <ol style="list-style-type: none"> <li>a) <input type="checkbox"/> Verified Statement Claiming Small Entity Status.</li> <li>b) <input type="checkbox"/> Copy of Notification of Missing Requirements.</li> <li>c) <input checked="" type="checkbox"/> Itemized receipt post card.</li> <li>d) <input checked="" type="checkbox"/> Serial number post card.</li> <li>e) <input checked="" type="checkbox"/> Written Opinion (PCT/IPEA/408) _____</li> <li>f) <input checked="" type="checkbox"/> Cover page of WO 01/00585 A1 _____</li> <li>g) <input type="checkbox"/> _____</li> <li>h) <input type="checkbox"/> _____</li> </ol> </li> </ol>		

U.S. Patent and Trademark Office  
 International Patent Cooperation Treaty  
 Patent Cooperation Treaty

<b>U.S. Application No.</b> <div style="font-size: 1.5em; font-weight: bold;">10/019229</div>	<b>International Application No.</b> PCT/BG00/02513	<b>Attorney Docket No.</b> 288748.0005
21. The following fees are submitted: <b>Basic National Fee (37 CFR 1.492(a)(1)-(5)):</b> <div style="margin-left: 20px;"> <input type="checkbox"/> Neither international preliminary examination fee nor international search fee paid to USPTO; International Search Report not prepared by the EPO or JPO \$ 1,040.00  <input checked="" type="checkbox"/> International preliminary examination fee not paid to USPTO; International Search Report prepared by EPO OR JPO \$ 890.00  <input type="checkbox"/> International preliminary examination fee not paid to USPTO; international search fee paid to USPTO \$ 740.00  <input type="checkbox"/> International preliminary examination fee paid to USPTO; all claims did not satisfy provisions of PCT Article 33(1)-(4) \$ 710.00  <input type="checkbox"/> International preliminary examination fee paid to USPTO; all claims satisfied provisions of PCT Article 33(1)-(4) \$ 100.00         </div>		<b>Calculations</b> <i>PTO USE ONLY</i>
<b>ENTER APPROPRIATE BASIC FEE AMOUNT =</b>		\$890.00
Surcharge of \$130.00 for furnishing the oath or declaration later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date 37 CFR 1.492(e)).		\$0.00
CLAIMS	NUMBER FILED	NUMBER EXTRA
Total Claims	31 - 20 =	11
Indep. Claims	7 - 3 =	4
Multiple Dependent Claims (if applicable)		\$280.00
<b>TOTAL OF ABOVE CALCULATIONS =</b>		\$1,704.00
Reduction by 1/2 for filing by small entity, if applicable. Verified Small Entity statement must also be filed. (Note 37 CFR 1.9, 1.27, 1.28)		\$0.00
<b>SUBTOTAL =</b>		\$1,704.00
Processing fee of \$130.00 for furnishing the English translation later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(f)).		\$0.00
<b>TOTAL NATIONAL FEE =</b>		\$1,704.00
Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31).		\$0.00
<b>TOTAL FEES ENCLOSED =</b>		\$1,704.00
<input checked="" type="checkbox"/> A check in the amount of \$1,704.00 to cover the above fees is enclosed. <input type="checkbox"/> Please charge my Deposit Account No. ____ in the amount of \$ ____ to cover the above fees. A duplicate copy of this sheet is enclosed. <input checked="" type="checkbox"/> The Director is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No.50-1656. A duplicate copy of this sheet is enclosed		
<b>SEND ALL CORRESPONDENCE TO:</b>  <b>John W. Ryan</b> <b>WILMER, CUTLER &amp; PICKERING</b> <b>2445 M. Street, N.W.</b> <b>Washington, D.C. 20037-1420</b>		<div style="text-align: right;">         John W. Ryan, Reg. No. 33,771        (202) 663-6446        (202) 663-6363 (facsimile)     </div> <div style="text-align: right; margin-top: 10px;"> <i>28 December 2003</i> </div>

JC13 Rec'd PCT/PTO 28 DEC 2001

PATENT APPLICATION  
IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Patent Application of

Enok TJØTTA et al.

Application No: [Not yet assigned]

Attn: **BOX PATENT APPLICATION**

Filed: December 28, 2001

Title: PYRAZOLIDINOL COMPOUNDS  
(National Stage Application of PCT/GB00/02513)

**PRELIMINARY AMENDMENT**

Commissioner of Patents  
Washington, D.C. 20231

Sir:

Prior to the examination of the above application, please amend this application as follows:

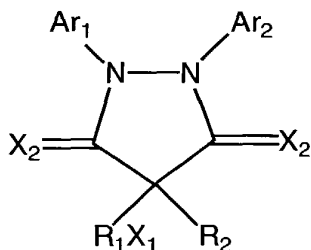
**IN THE ABSTRACT OF THE DISCLOSURE:**

Please employ the attached separate page to this amendment as the Abstract of the Disclosure.

**IN THE CLAIMS:**

Please AMEND the claims as follows:

1. (Amended) A compound of formula I



(I)

## PATENT APPLICATION

(where each  $X_2$ , which may be the same or different is O or S,

$X_1$  is O, OO or S,

$R_1$  is hydrogen or a hydroxyl or thiol protecting group,

$R_2$  is hydrogen or a alkyl, alkenyl, alkynyl, alkaryl, aralkyl or aralkenyl group, containing up to 10 carbons, optionally substituted by a sulphonyl group,

and each of

$Ar_1$  and  $Ar_2$ , which may be the same or different, is a homo or heterocyclic aromatic group comprising 5 to 7 membered aromatic ring, optionally carrying a fused aromatic ring and optionally substituted on ring atoms by  $C_{1-6}$  alkyl, hydroxy, thiol,  $C_{1-6}$  alkoxy, cyano, Cl, F, Br, I, protected hydroxy, or protected thiol), or a physiologically acceptable salt thereof, for the manufacture of a medicament for use in therapy or prophylaxis.

3. (Amended) The method as claimed in claim 2 comprising administering said compound, or a physiologically acceptable salt thereof in combination with another antiviral agent.

4. (Amended) The method as claimed in claim 3 wherein said additional antiviral agent is at least one antiviral agent selected from a reverse transcriptase inhibitor and a protease inhibitor.

5. (Amended) The method as claimed in claim 3 wherein said additional antiviral agent is an agent selected from the group of AZT, indinavir, nevirapine and 2',3'-dideoxyinosine (ddI).

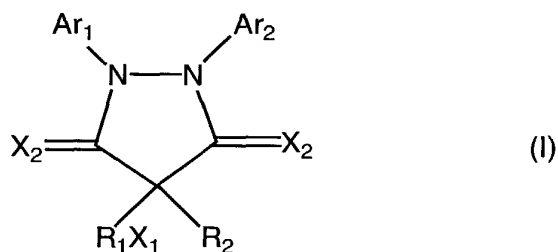
6. (Amended) The method as claimed in any of claims 2 to 5 wherein said disease is a disease caused by a pathogen from the group of togaviridea, reoviridea, picornaviridea, hantaviridea, orthomyxoviridea, paramyxoviridea, mononegaviralis, viral hepatitis, haemorrhagic fevers, flaviviridea, viral encephalitis, coronaviridea, calciviridea, adenoviridea, papovaviridea, arboviridea, pox virus, rhabdoviridea, arenaviridea HIV-1, HIV-2, HTLV-I, HTLV-II and herpes viruses.

PATENT APPLICATION

8. (Amended) The method of combatting HIV infection as claimed in claim 7 wherein said T-lymphocyte growth suppressing agent is pyrazolidinol.

9. (Amended) The method as claimed in claim 8 wherein said interval is at least 9 months.

10. (Amended) The method as claimed in claim 9 wherein a compound of formula I



(where each  $X_2$ , which may be the same or different is O or S,

$X_1$  is O, OO or S,

$R_1$  is hydrogen or a hydroxyl or thiol protecting group,

$R_2$  is hydrogen or a alkyl, alkenyl, alkynyl, alkaryl, aralkyl or aralkenyl group, containing up to 10 carbons, optionally substituted by a sulphonyl group, and each of  $Ar_1$  and  $Ar_2$ , which may be the same or different, is a homo or heterocyclic aromatic group comprising 5 to 7 membered aromatic ring, optionally carrying a fused aromatic ring and optionally substituted on ring atoms by  $C_{1-6}$  alkyl, hydroxy, thiol,  $C_{1-6}$  alkoxy, cyano, Cl, F, Br, I, protected hydroxy, or protected thiol), or a physiologically acceptable salt there is administered in a daily dose of 0.1 to 10  $\mu\text{mol/kg}$  bodyweight.

12. (Amended) The pharmaceutical composition as claimed in claim 11 additionally comprising another antiviral agent.

14. (Amended) The compound as claimed in claim 13 wherein one  $X_2$  group is S.

15. (Amended) The compound as claimed in claim 13 wherein  $X_1$  is O.

## PATENT APPLICATION

16. (Amended) The compound as claimed in claim 13 wherein  $R_1$  is acyl.
17. (Amended) The compound as claimed in claim 13 wherein  $R_1$  is hydrogen.
18. (Amended) The compound as claimed in claim 13 wherein each  $X_2$  is oxygen,  $R_1X_1$  is HO or  $CH_3CO.O$ , and  $R_2$  is  $C_{1-6}$  alkyl or alkenyl, or a salt thereof.
19. (Amended) The compound as claimed in any of claims 13 to 18 for use as a medicament.
20. (Amended) A compound comprising 4-butyl-4-hydroxy-2(p-hydroxyphenyl)-1-phenyl-3,5-pyrazolidinedione for use as a medicament.
22. (Amended) The method of claim 21 wherein said disease is selected from Addison's disease, Behçet's syndrome, diabetes mellitus, haemolytic anaemia, lupus erythematosus, multiple sclerosis, myasthenia gravis, pernicious anaemia, polyglandular deficiency, polymyositis, dermatomyositis, testicular failure, thrombocytopenic purpura, Chrons disease, ulcerative colitis and rheumatoid arthritis.
23. (Amended) The method of claim 21 wherein said tissue rejection is tissue rejection following transplant.

PATENT APPLICATION

REMARKS

Claims 1, 9, 10, 14-17 and 20 have been amended.

The Examiner is respectfully requested to consider this preliminary amendment prior to examination of the application. The above amendments have been made to place the claims in conformance with U.S. practice. The applicant has reviewed and amended the claims for clarification purposes only. There has been no narrowing amendments entered and no new matter has been added.

Attached as an appendix entitled "**Version with Markings to Show Changes Made**" is a marked-up version of the presently amended claims.

Please charge any fees due in connection with this request to undersigned's Deposit Account No. 50-1656.

Respectfully submitted,

WILMER CUTLER & PICKERING

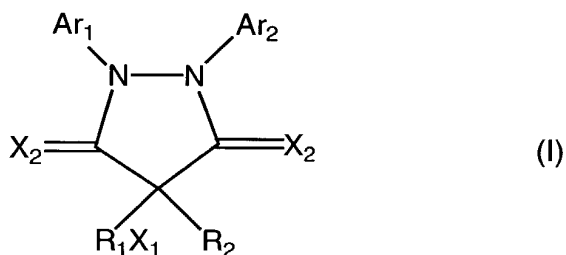
Date: December 28, 2001

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1. [The use of a] A compound of formula I



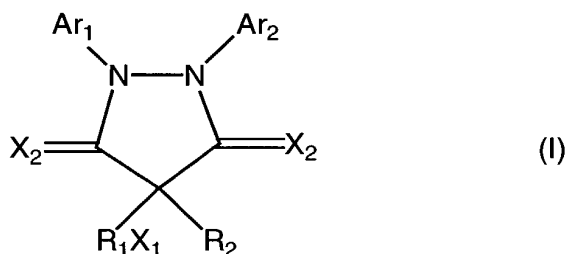
Ar<sub>1</sub> and Ar<sub>2</sub>, which may be the same or different, is a homo or heterocyclic aromatic group comprising 5 to 7 membered aromatic ring, optionally carrying a fused aromatic ring and optionally substituted on ring atoms by C<sub>1-6</sub> alkyl, hydroxy, thiol, C<sub>1-6</sub> alkoxy, cyano, Cl, F, Br, I, protected hydroxy, or protected thiol), or a physiologically acceptable salt thereof, for the manufacture of a medicament for use in therapy or prophylaxis.

5. [A] The method as claimed in claim 3 wherein said additional antiviral agent is an agent selected from the group of AZT, indinavir, nevirapine and 2',3'-dideoxyinosine (ddI).



## PATENT APPLICATION

6. [A] The method as claimed in any of claims 2 to 5 wherein said disease is a disease caused by a pathogen from the group of togaviridea, reoviridea, picornaviridea, hantaviridea, orthomyxoviridea, paramyxoviridea, mononegaviralis, viral hepatitis, haemorrhagic fevers, flaviviridea, viral encephalitis, coronaviridea, calciviridea, adenoviridea, papovaviridea, arboviridea, pox virus, rhabdoviridea, arenaviridea HIV-1, HIV-2, HTLV-I, HTLV-II and herpes viruses.
8. [A] The method of combatting HIV infection as claimed in claim 7 wherein said T-lymphocyte growth suppressing agent is pyrazolidinol.
9. [A] The method as claimed in [claim 7 or] claim 8 wherein said interval is at least 9 months.
10. [A] The method as claimed in [any of claims 7 to] claim 9 wherein a compound of formula I



(where each X<sub>2</sub>, which may be the same or different is O or S,

X<sub>1</sub> is O, OO or S,

R<sub>1</sub> is hydrogen or a hydroxyl or thiol protecting group,

R<sub>2</sub> is hydrogen or an alkyl, alkenyl, alkynyl, alkaryl, aralkyl or aralkenyl group, containing up to 10 carbons, optionally substituted by a sulphonyl group, and each of Ar<sub>1</sub> and Ar<sub>2</sub>, which may be the same or different, is a homo or heterocyclic aromatic group comprising 5 to 7 membered aromatic ring, optionally carrying a fused aromatic ring and optionally substituted on ring atoms by C<sub>1-6</sub> alkyl, hydroxy, thiol, C<sub>1-6</sub> alkoxy, cyano, Cl, F, Br, I, protected hydroxy, or protected thiol), or a physiologically acceptable salt there is administered in a daily dose of 0.1 to 10 μmol/kg bodyweight.

## PATENT APPLICATION

12. [A] The pharmaceutical composition as claimed in claim 11 additionally comprising another antiviral agent.
14. [A] The compound as claimed in claim 13 [or claim 14] wherein one  $X_2$  group is S.
15. [A] The compound as claimed in [either of claims] claim 13 [or 14] wherein  $X_1$  is O.
16. [A] The compound as claimed in [any of claims] claim 13 [to 15] wherein  $R_1$  is acyl.
17. [A] The compound as claimed in [any of claims] claim 13 [to 16] wherein  $R_1$  is hydrogen.
18. [A] The compound as claimed in claim 13 wherein each  $X_2$  is oxygen,  $R_1X_1$  is HO or  $CH_3CO.O$ , and  $R_2$  is  $C_{1-6}$  alkyl or alkenyl, or a salt thereof.
19. [A] The compound as claimed in any of claims 13 to 18 for use as a medicament.
20. A compound comprising 4-butyl-4-hydroxy-2(p-hydroxyphenyl)-1-phenyl-3,5-pyrazolidinedione for use as a medicament.
22. [A] The method of claim 21 wherein said disease is selected from Addison's disease, Behçet's syndrome, diabetes mellitus, haemolytic anaemia, lupus erythematosus, multiple sclerosis, myasthenia gravis, pernicious anaemia, polyglandular deficiency, polymyositis, dermatomyositis, testicular failure, thrombocytopenic purpura, Chrons disease, ulcerative colitis and rheumatoid arthritis.
23. [A] The method of claim 21 wherein said tissue rejection is tissue rejection following transplant.

## PATENT APPLICATION

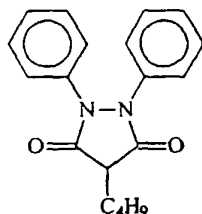
### ABSTRACT OF THE DISCLOSURE

The invention provides the use of an optionally hydroxy-protected 4-hydroxyo or hydroperoxy-3,5-dioxopyrazolidine or an equivalent wherein a pyrazolidine ring attached oxygen is replaced by a sulphur, or a physiologically acceptable salt thereof, for the manufacture of a medicament for use in therapy or prophylaxis. Additionally, the invention provides a method of combating HIV infection which comprises administering to an HIV-infected patient a T-lymphocyte growth suppressing agent, preferably a pyrazolidinol, in an amount sufficient to suppress T-lymphocyte growth in said patient for a period sufficient to reduce the T-lymphocyte concentration in lymph nodes in said patient by at least 25%, said administration being repeated at intervals of at least 3 months.

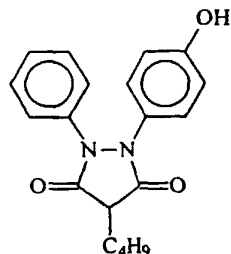
Pyrazolidinol Compounds

The present invention relates to certain pyrazolidinols and their sulphur (i.e. thio/thiol) analogs and pharmaceutical compositions thereof for use in antiviral, e.g. anti-HIV therapy and as anti-inflammatories and immunomodulators.

Phenbutazone and oxyphenbutazone are 1,2-bis aromatic-3,5-pyrazolidinediones which have been used as non-steroidal anti-inflammatory drugs (NSAIDs)



Phenbutazone (PB)



Oxyphenbutazone (OPB)

Other 3,5-pyrazolidinediones have likewise been proposed for use as NSAIDs (see for example US-A-3968219 (Rahtz)) and the hydroxy-protected enol forms have been proposed as pro-drug forms of phenbutazone and oxyphenbutazone (see US-A-4117232 (Bodor), US-A-3957803 (Bodor), US-A-4169147 (Bodor), US-A-4036845 (Bodor) and US-A-4139709 (Bodor)).

In US-A-4956377 (Miesch) it was proposed that this class of NSAIDs had utility as an antiviral agent, in particular for the treatment of HIV.

We have now surprisingly found that where the 4-carbon of the N<sub>2</sub>C<sub>3</sub> ring carries an optionally protected

hydroxy or thiol group, the compounds have very significantly enhanced antiviral, in particular anti-HIV, efficacy.

Thus viewed from one aspect the invention provides the use of an optionally hydroxy-protected 4-hydroxy or hydroperoxy-3,5-dioxo-pyrazolidine or an equivalent wherein a pyrazolidine ring attached oxygen is replaced by a sulphur, or a physiologically acceptable salt thereof, for the manufacture of a medicament for use in therapy or prophylaxis.

Where a particular 4-hydroxy or hydroperoxy-3,5-dioxo-pyrazolidine may exist in more than one stereoisomeric form, it may be used in single isomer form or as an isomer mixture, e.g. a racemic mixture.

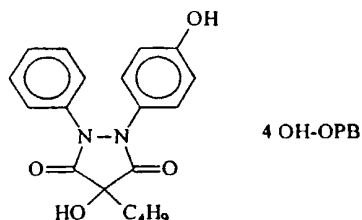
Viewed from a further aspect, the invention provides an optionally hydroxy-protected 4-hydroxy or hydroperoxy-3,5-dioxo-pyrazolidine or an equivalent wherein a pyrazolidine ring attached oxygen is replaced by a sulphur, or a physiologically acceptable salt thereof.

Viewed from a still further aspect the invention provides a method of treatment of the human or non-human (e.g. mammalian, reptilian or avian) body to combat an inflammatory or viral disease, preferably an immunodeficiency viral disease, in particular HIV, which method comprises administering to said body an optionally hydroxy-protected 4-hydroxy or hydroperoxy-3,5-dioxo-pyrazolidine or an equivalent wherein a pyrazolidine ring attached oxygen is replaced by a sulphur, or a physiologically acceptable salt thereof.

Viewed from a still further aspect, the invention provides a pharmaceutical composition comprising an optionally hydroxy-protected 4-hydroxy or hydroperoxy-3,5-dioxo-pyrazolidine or an equivalent wherein a pyrazolidine ring attached oxygen is replaced by a sulphur, or a physiologically acceptable salt thereof, together with at least one pharmaceutically acceptable

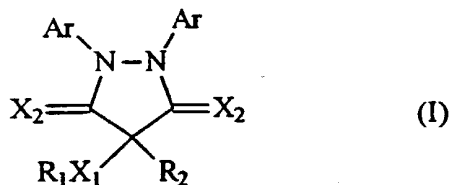
carrier or excipient.

The applicants have found that oxyphenbutazone, as commercially available, contains minute quantities of certain impurities, presumably as a result of undesired oxidative breakdown. One of these, present at about 0.4% wt, is 4-butyl-4-hydroxy-2(p-hydroxyphenyl)-1-phenyl-3,5-pyrazolidinedione (hereinafter "4-OH-OPB"), i.e.



4-OH-OPB is of course a compound according to the invention and thus it should be understood that references to the 4-hydroxy compounds of the invention, their use and compositions thereof should not be taken to include references to such compounds when in intimate admixture with overwhelmingly larger quantities of a 3,5-pyrazolidinedione which carries no optionally protected 4-hydroxy or 4-thiol group. By overwhelmingly larger is meant a relative weight ratio of at least 98:2. In general, the compounds of the invention should not desirably be used in intimate admixture with larger quantities (i.e. a relative weight ratio of more than 50:50) of such compounds carrying no O or S attached group at the 4-position, and more desirably they should not be used with such compounds present in greater than 10:90 weight ratio.

The compounds of the invention, hereinafter referred to as pyrazolidinols for convenience, will preferably be of formula I



- 4 -

(where each  $X_2$ , which may be the same or different is O or S, preferably O;

$X_1$  is O, OO or S, preferably O or S, most preferably O;

$R_1$  is hydrogen or a hydroxyl or thiol protecting group (e.g. an acyl group, preferably containing up to 6 carbons, e.g. an acyl group such as an alkylcarbonyl group, for example acetyl), preferably hydrogen;

$R_2$  is hydrogen or more preferably a carbon attached organic group containing up to 10 carbons, e.g. an alkyl, alkenyl, alkynyl, alkaryl, aralkyl or aralkenyl group, optionally substituted, e.g. by a sulphonyl group; and

each Ar, which may be the same or different, is a homo or heterocyclic aromatic group, optionally substituted, e.g. by  $C_{1-6}$  alkyl or alkoxy groups) or a salt thereof.

In the compounds of the invention 0, 1 or 2 of the  $X_1$  and  $X_2$  groups may be S. It is thought that it is especially preferred that one thio  $X_2$  group be present.

In the compounds of the invention, the  $R_2$  group is preferably other than hydrogen and may for example be straight chain, branched, cyclic or cyclic-attached-to-straight chain. Preferably it is an alkyl or alkenyl group, especially a  $C_{1-6}$  alkyl or alkenyl group, e.g. n-propyl, n-butyl, n-pentyl or 1-methyl-but-2-en-4-yl or an aralkyl (e.g. benzyl) or alkaryl (e.g. methylphenyl) or arylsulphonylalkyl (eg phenylsulphonylethyl) group.

Where  $R_1$  in the compounds of the invention is other than hydrogen it is preferably a metabolically labile hydroxy- or thiol-protecting group which yields a physiologically tolerable  $R_1OH$  metabolite. Acyl groups are preferred in this regard.

In the compounds of formula I, where each  $X_2$  is oxygen and one Ar is phenyl, the other Ar is preferably other than phenyl e.g. parahydroxyphenyl.

A wide range of hydroxy- and thiol-protecting groups however is known from the literature (see McOmie, "Protective groups in organic chemistry", Plenum, 1973

- 5 -

and Greene, "Protective groups in organic synthesis", Wiley Interscience, NY, 1981) and many compounds of formula I in which  $R_1$  is a protecting group may be useful as intermediates in the production of compounds of formula I in which  $R_1$  is hydrogen.

The Ar groups in the compounds of formula I are preferably 5 to 7 membered aromatic rings, optionally carrying a fused aromatic ring and optionally substituted on ring atoms, for example by  $C_{1-6}$  alkyl groups but especially by electron withdrawing substituents, e.g. hydroxy, thiol, phenyl,  $C_{1-6}$  alkoxy, cyano, halo (e.g. Cl, F, Br or I), protected hydroxy, or protected thiol. Ring heteroatoms will generally be selected from O, N and S, preferably with a single ring heteroatom in any aromatic Ar heterocycle. Ar is preferably phenyl optionally substituted, especially in the para-position by  $-X_1-R_1$  or Cl (where  $-X_1-R_1$  is as defined above). Especially preferably one Ar is phenyl and the other is p-hydroxy-phenyl.

Where the substitution of the pyrazolidinols of the invention is such that they may form addition salts with acids or bases, the addition salts which have physiologically tolerable counterions are of course preferred, e.g. sodium, organic amine, halides, phosphates, hydrogen carbonates, etc.

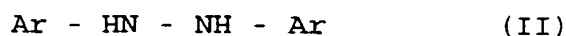
The pyrazolidinols of the invention may particularly advantageously be used in combination therapy with other antiviral, especially anti-HIV, agents, in particular reverse transcriptase inhibitors and/or protease inhibitors, e.g. zidovudine, didanosine, zalcitabine, stavudine, lamivudine, nevirapine, delavirdine, indinavir, ritonavir, nelfinavir, hydroxyurea, colchicine, AZT and 2',3'-dideoxyinosine (ddI). Such combination therapy forms a further aspect of the present invention.

A drawback of traditional combination therapy, has often been that even under intensive antiviral treatment with a combination of drugs, a little HIV production

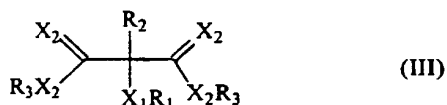


continues and is unaffected by treatment. The compounds of the invention may prove to have an effect in reducing this residual HIV production when given in combination with other antiviral agents. This may be due to the increasing antiviral effect which has been seen in long term cell culture experiments and which may counteract any development of resistance to the compounds.

The pyrazolidinols of the invention may be prepared by oxidation of a corresponding compound where  $R_1X_1$  is replaced by hydrogen; by reaction of a corresponding compound where  $R_1X_1$  is  $HX_1$  with a hydroxy or thiol protecting agent to introduce a non-hydrogen  $R_1$  group; or by condensation of a hydrazine derivative with an optionally protected 2-hydroxy-propane dioic acid ester (or a sulphur analog), e.g. by condensation of a compound of formula II

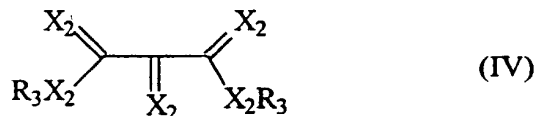


with a compound of formula III



where  $R_1$ ,  $R_2$ ,  $X_1$ ,  $X_2$  and Ar are as hereinbefore defined and  $X_2R_3$  is a leaving group, for example where  $R_3$  is an alkyl group, e.g. a  $C_{1-6}$  alkyl group.

Alternatively, a compound of formula II may be condensed with a compound of formula IV



(where  $X_2$  and  $R_3$  are as defined above) and then reacted with an alkylating agent, e.g.  $(R_2)_2\text{Zn}$  to produce a compound of formula I in which  $X_1R_1$  is OH or SH.

- 7 -

For administration, the pyrazolidinols of the invention may be formulated in any convenient form, e.g. tablets, coated tablets (e.g. delayed release tablets), capsules, solutions, suspensions, dispersions, syrups, powders, sprays, suppositories, transdermal patches, gels, emulsions and creams. Administration may be via any convenient route, e.g. oral, rectal, transdermal, nasal, subcutaneous, intravenous, intramuscular, etc. Oral administration, e.g. of tablets or capsules is preferred. The pyrazolidinols may be formulated together with conventional pharmaceutical carriers, diluents or excipients, e.g. aqueous carriers (for example water for injections), binders, fillers, stabilizers, osmolality adjusting agents, effervescing agents, pH modifiers, viscosity modifiers, sweeteners, lubricants, emulsifiers, flavours, coating agents (e.g. gastric juice resistant coatings), etc. Where any formulation results in a loss of compound, this loss should be calculated and the dosage increased proportionally to obtain the desired active concentration.

The dosage of the pyrazolidinols given according to the invention will depend on the size and species of the subject being treated but will generally be in the range of 0.05 to 2000 mg/day, more particularly 0.5 to 1000 mg/day, especially 1 to 100 mg/day, preferably with administration being effected once, twice, three times or four times daily. For mice, doses of up to 2000 mg/kg (corresponding to 20 mM maximal concentration in extracellular fluid) could be given before lethal dosage was reached, ie effective treatment doses were up to 200000 times smaller than the lethal dose.

For regular, e.g. continuous daily treatment according to the invention, the daily dosage of the pyrazolidinol will preferably be in the range 5 nmol to 2  $\mu$ mol/kg bodyweight, more preferably 100 nmol to 1.5  $\mu$ mol/kg, especially 500 nmol to 1  $\mu$ mol/kg.

Inhibition of virus production may be achieved by small intermittent doses of pyrazolidinol, and are expected to induce inhibition of the virus after a latency of about 11 weeks. Subsequently, inhibition may be expected to level out, should resistance to the compound develop. Such doses may be administered at a frequency of 1-14 days, preferably 7 days. The doses should be equivalent to a concentration in plasma/tissue fluid of from 100-1000 nM and may be obtained by ingestion or injection of from 0.7-7 mg in a 70 Kg human.

However, in a particularly preferred embodiment of the invention, a pyrazolidinol according to the invention is administered at a dose sufficient to suppress T-lymphocyte (CD4 and CD8 cell) growth (e.g. a daily dose of 0.1 to 10  $\mu$ mol/kg) for a period of 1 to 14 days, preferably 2 to 7 days at intervals of at least 3 months, preferably at least 9 months, e.g. 10 to 18 months. In this way the patient's immune system may be "refreshed" by removal of the preponderance of T-lymphocytes directed to HIV antigens. Such a treatment indeed is novel and forms a further aspect of the invention. Viewed from this aspect the invention provides a method of combatting HIV infection which comprises administering to an HIV-infected patient a T-lymphocyte growth suppressing agent, e.g. a pyrazolidinol, in an amount sufficient to suppress T-lymphocyte growth in said patient for a period sufficient to reduce the T-lymphocyte concentration in the lymphatic system, e.g. the lymph nodes, in said patient by at least 25%, more preferably at least 50%, said administration being repeated at intervals of at least 3 months, preferably at least 9 months.

High tissue concentrations intended to give an immunomodulating effect should preferably be given for limited periods at doses of 1 $\mu$ M or above in plasma/tissue fluid. Such doses and lengths of

- 9 -

administration will vary according to the condition of each patient and may be decided with the guidance of tests such as the count of HIV memory subsets of T8 and T4. As stated above, the goal of treatment according to this aspect of the invention should be to reduce subsets which are found to be too prevalent without overly affecting naive T-cells.

In order to obtain the desired reduction of HIV specific lymphocytes (e.g. HIV memory CD8 and CD4 lymphocytes) without overly affecting naive T-lymphocytes or other essential blood cells, monoclonal antibodies against the unwanted subtypes may also be administered. Further, drugs such as kolchicine and/or hydroxy-urea may be included in the intermittent intensive treatment. Such additional drugs are anticipated to have a somewhat different immunomodulating effect to the compounds of the invention and so may be used advantageously in combination with pyrazolidinols for refreshing the immune system.

Besides HIV, the pyrazolidinols of the invention may be used to combat other viral infections, especially retroviral infections but also infections by togaviridea, reoviridea, picornaviridea, hantaviridea, orthomyxoviridea, paramyxoviridea, mononegaviralis, viral hepatitis, haemorrhagic fevers, flaviviridea, viral encephalitis, coronaviridea, calciviridea, adenoviridea, papovaviridea, arboviridea, pox virus, rhabdoviridea, herpes virus and arenaviridea. The pyrazolidinols of the invention may in particular be used to combat viral infections of CD4 cells, e.g. HIV-1, HIV-2, HTLV-I, HTLV-II and herpes viruses, for example to combat AIDS, T-cell tumours (e.g. Sezary Syndrome, mycosis fungoides and T-cell lymphoma, and particularly CD4 cell tumours), tropic spastic paraparesis, and Kaposi's sarcoma. Moreover despite not being of the accepted formula for NSAIDs (which

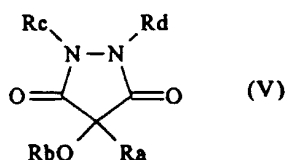
- 10 -

would require an acid proton in place of  $R_1X_1$  at the 4-position), they may be used as anti-inflammatory drugs. All these uses form aspects of the invention.

Due to the immunomodulating effect of the compounds of the invention, they are expected to have uses in control of other immune-system related diseases, such as auto immune diseases and as immunosuppressants. In particular, the compounds of the invention are expected to have a positive effect on the generation of autoimmune diseases, on developed autoimmune diseases and on diseases related to such diseases, such as Addison's disease, Behçet's syndrome, diabetes mellitus and other endocrine diseases, haemolytic anaemia, lupus erythematosus, multiple sclerosis, myasthenia gravis, pernicious anaemia, polyglandular deficiency, polymyositis, dermatomyositis, testicular failure, thrombocytopenic purpura, Crohns disease, ulcerative colitis, rheumatic disorders (e.g. rheumatoid arthritis) etc.

The effect of the compounds of the invention on the immune system may also be that of immunosuppression. Such an effect may be used, for example, to control rejection of a medical transplant or implant. In particular, the compounds may be used to reduce rejection following tissue or organ transplant.

Various 4-hydroxy-3,5-dioxo-pyrazolidines are known in the literature (although not for medical purposes such as HIV therapy). These are compounds of formula V



where  $R_a$  to  $R_d$  are as set out in Table 1 below:

Table 1

$R_a$	$R_b$	$R_c$	$R_d$
H	H	H	$C_6H_5$
H	H	$C_6H_5$	$C_6H_5$
$CH_3$	H	H	$C_6H_5$
$CH_3$	H	H	$-CH_2-C_6H_5$
$CH_3$	H	H	$p-CH_3O-C_6H_4$
$CH_3$	H	H	$p-Cl-C_6H_4$
$C_2H_5$	H	H	$C_6H_5$
$C_2H_5$	H	$C_6H_5$	$C_6H_5$
$C_2H_5$	H	H	N-methyl-piperidin-4-yl
$iC_3H_7$	H	H	$C_6H_5$
$nC_3H_7$	H	H	$C_6H_5$
$nC_3H_7$	H	$C_6H_5$	$C_6H_5$
$nC_3H_7$	H	H	5-phenyl-triazol-1-yl
$C_4H_9$	H	H	$C_6H_5$
$C_4H_9$	H	$C_6H_5$	$C_6H_5$
$C_4H_9$	H	$C_6H_5$	$p-OH-C_6H_4$
$C_4H_9$	OH	$C_6H_5$	$C_6H_5$
$C_4H_9$	OH	$C_6H_5$	$p-OH-C_6H_4$
$C_4H_9$	H	H	N-methyl-piperidin-4-yl
$C_5H_{11}$	H	H	$C_6H_5$
$C_5H_{11}$	H	$C_6H_5$	$C_6H_5$
$C_5H_{11}$	H	H	5-phenyl-triazol-1-yl
Cyclohexyl	H	H	$C_6H_5$
Phenyl	H	H	$C_6H_5$
Phenyl	H	$C_6H_5$	$C_6H_5$
Benzyl	H	H	$C_6H_5$
Benzyl	H	$C_6H_5$	$C_6H_5$
$CH_3CO(CH_2)_2$	H	$C_6H_5$	$C_6H_5$
$(CH_3)_2C=CH-$	H	$C_6H_5$	$C_6H_5$
$(CH_2)_2C=CHCH_2$	H	$C_6H_5$	$C_6H_5$
$C_6H_5SCH_2CH_2$	H	$C_6H_5$	$C_6H_5$
Pyrrolidin-1-yl	H	$C_6H_5$	$C_6H_5$
Piperidin-1-yl	H	$C_6H_5$	$C_6H_5$
Morpholin-4-yl	H	$C_6H_5$	$C_6H_5$

Such compounds are thus not claimed per se herein; however their use and pharmaceutical compositions containing them do form part of the scope of the invention.

The invention will now be illustrated further by the following non-limiting Examples and by reference to the Figures, in which:

Figure 1 shows the HIV antigen concentration in human CD4 cells infected with HIV and treated with 4-butyl-4-hydroxy-2(p-hydroxyphenyl)-1-phenyl-3,5-pyrazolidinedione (4OH-OPB) or phenbutazone (PB) at various concentrations;

Figure 2 shows the effect of 4-butyl-4-hydroxy-2(p-hydroxyphenyl)-1-phenyl-3,5-pyrazolidinedione (4OH-OPB) when used in combination with AZT;

Figure 3 shows the effect of 4-butyl-4-hydroxy-2(p-hydroxyphenyl)-1-phenyl-3,5-pyrazolidinedione (4OH-OPB) when used in combination with indinavir; and;

Figure 4 shows the effect of 4-butyl-4-hydroxy-2(p-hydroxyphenyl)-1-phenyl-3,5-pyrazolidinedione (4OH-OPB) when used in combination with nevirapine;

A similar effect to those shown in figures 2-4 is seen when OPB is used in combination with 2',3'-dideoxyinosine (ddI).

#### **EXAMPLE 1**

##### **Preparation of 4-Methoxyazobenzene**

A mixture of 4-phenylazophenol (9.9g; 50 mmol), iodomethane (7.1 g; 50 mmol), potassium carbonate (6.9 g; 50 mmol), and acetone (100 ml) was refluxed 48 h. After evaporating off the solvent, the residue was dissolved in water (25 ml), diethyl ether (50 ml) and THF (30 ml). The aqueous layer was extracted with ether (3 x 20 ml) and the combined organic solutions were washed with saturated NaCl solution (1 x 20 ml) and

- 13 -

dried ( $\text{MgSO}_4$ ). After filtration and evaporation, the residue was recrystallized from 96% ethanol to give 8.7 g (82%).

#### EXAMPLE 2

##### Preparation of 1-(4-Methoxyphenyl)-2-phenylhydrazine

Zinc powder (10.0 g; 0.15 mol) was added to a stirred mixture of 4-methoxyazobenzene (4.24 g; 20.0 mmol) in 96% ethanol (75 ml) and saturated  $\text{NH}_4\text{Cl}$  solution (2.0 ml) at 0 °C (bath temperature). Two more portions of saturated  $\text{NH}_4\text{Cl}$  solution (2.0 ml) were added at 1.5 h intervals. The yellowish solution was poured into cold water (100 ml) and filtered. The residue was extracted with methylene chloride (5 × 50 ml). The combined aqueous phases were extracted with methylene chloride (3 × 25 ml). The combined organic solutions were dried ( $\text{Na}_2\text{SO}_4$ ), filtered, and evaporated to give 4.3 g crude 1-(4-methoxyphenyl)-2-phenylhydrazine as a reddish oil.

#### EXAMPLE 3

##### Preparation of 4-(1-Butyl)-1-(4-methoxyphenyl)-2-phenyl-3,5-pyrazolidinedione

Diethyl butylmalonate (4.33 g; 20.0 mmol) was added to a stirred solution of sodium (0.46 g; 20.0 mmol) in absolute ethanol (20 ml), followed by crude 1-(4-methoxyphenyl)-2-phenylhydrazine (4.3 g; 20 mmol max.) in absolute ethanol (5 ml). About 2/3 of the ethanol was distilled off and xylene (20 ml) was added to the residue. The reaction mixture was heated to 140-145 °C (bath temperature) for 15 h to distill off the rest of the ethanol. The reaction mixture was cooled to 0 °C (bath temperature) and poured into ice water (ca. 100 ml). The aqueous layer was extracted with  $\text{CH}_2\text{Cl}_2$  (2 × 15



ml); the extracts were discarded. The cold aqueous layer was acidified with 6 M HCl (5 ml) and extracted with  $\text{CH}_2\text{Cl}_2$  (3 × 10 ml). The combined extracts were washed with water (2 × 10 ml) and dried ( $\text{MgSO}_4$ ). Filtration and evaporation gave 3.84 g amber oil. Purified by flash chromatography on a 130 × 65 mm silica gel 60 column eluted with ethyl acetate-heptane (1:3) to give 1.45 g (21%) colourless oil.

$^1\text{H}$  NMR (200 MHz;  $\text{CDCl}_3$ ):  $\delta$  0.90 (3H, t,  $J$  = 7.5 Hz), 1.25-1.6 (4H, m), 2.0-2.15 (2H, m), 3.37 (3H, t,  $J$  = 6.0 Hz), 3.69 (3H, s), 6.81 (2H, d,  $J$  = 8.4 Hz), 7.22 (2H, d,  $J$  = 8.6 Hz), 7.1-7.35 (5H, m).

$^{13}\text{C}$  NMR (50 MHz;  $\text{CDCl}_3$ ):  $\delta$  13.6, 22.2, 27.5, 27.6, 45.6, 54.7, 112.9, 121.6, 123.3, 125.4, 127.1, 127.4, 133.9, 156.5, 168.2, 168.7.

#### EXAMPLE 4

##### Preparation of 1,2-Diphenyl-4-(4-methylphenyl)-3,5-pyrazolidinedione

Prepared from 1,2-diphenylhydrazine (3.70 g; 20.0 mmol), diethyl 2-(p-tolyl)malonate (5.0 g; 20.0 mmol), and sodium (0.46 g; 20.0 mmol) using the procedure of Example 3. The crude product crystallized on standing and was recrystallized twice from absolute ethanol to give 1.22 g (18%), mp 184-185 °C.

$^1\text{H}$  NMR (200 MHz;  $\text{CDCl}_3$ ):  $\delta$  2.31 (3H, s), 4.51 (1H, s), 7.1-7.4 (14H, m).

$^{13}\text{C}$  NMR (50 MHz;  $\text{CDCl}_3$ ):  $\delta$  21.1, 51.9, 122.7, 126.9, 128.3, 129.0, 129.9, 135.8, 138.3, 168.6.

#### EXAMPLE 5

- 15 -

Preparation of 4-Benzyl-1,2-diphenyl-3,5-pyrazolidinedione

Prepared from 1,2-diphenylhydrazine (4.60 g; 25.0 mmol), diethyl benzylmalonate (5.0 g; 20 mmol), and sodium (0.46 g; 20.0 mmol) using the procedure of Example 3. The crude product was recrystallized from absolute ethanol to give 3.51 g (50%), mp 136-137 °C [lit. 137-138 °C (Beil. III/IV, **24**, 1463)].

<sup>1</sup>H NMR (200 MHz; CDCl<sub>3</sub>): δ 3.41 (2H, d, J = 4.6 Hz), 3.63 (1H, t, J = 5.0 Hz), 6.85-7.3 (10H, m).

<sup>13</sup>C NMR (50 MHz; CDCl<sub>3</sub>): δ 33.9, 48.5, 123.2, 126.9, 127.3, 128.6, 128.7, 129.9, 135.2, 135.4, 169.3.

EXAMPLE 6

Preparation of 4-Allyl-1,2-diphenyl-3,5-pyrazolidinedione

Prepared from 1,2-diphenylhydrazine (5.2 g; 28.0 mmol), diethyl allylmalonate (5.0 g; 25.0 mmol), and sodium (0.58 g; 25.0 mmol) using the procedure of Example 3. The crude product was recrystallized from absolute ethanol to give 2.21 g (30%) tan crystals, mp 135-137 °C.

<sup>1</sup>H NMR (200 MHz; CDCl<sub>3</sub>): δ 2.82 (2H, t, J = 6.0 Hz), 3.46 (2H, t, J = 5.4 Hz), 5.1-5.3 (2H, dd), 5.7-5.95 (1H, m), 7.1-7.3 (10H, m).

<sup>13</sup>C NMR (50 MHz; CDCl<sub>3</sub>): δ 31.7, 46.4, 119.9, 122.7, 126.8, 128.9, 131.7, 135.6, 169.5.

EXAMPLE 7

Preparation of 4-(1-Butyl)-4-hydroxy-1-(4-hydroxyphenyl)-2-phenyl-3,5-pyrazolidinedione (4OH-OPB)

**Method A**

Oxyphenbutazone.H<sub>2</sub>O (1 mmol), 30% H<sub>2</sub>O<sub>2</sub> (0.7 mL), 1N NaOH (0.1 mL) and methanol (3.5 mL) are allowed to stand for 13 hours at ambient temperature. The mixture is then poured into 5% HCl (20 mL) and extracted with ethyl acetate (2 x 20 mL). The ethyl acetate phase is separated, dried over sodium carbonate and the solvent is removed under reduced pressure without heating. The residue is subjected to flash chromatography (silica/ethyl acetate). The title product is recrystallized from ethyl acetate.

**Method B**

A solution of oxyphenbutazone hydrate (2.0 g; 5.8 mmol), 35% hydrogen peroxide solution (3.4 ml; 40 mmol), and 1 M sodium hydroxide solution (0.6 ml; 0.6 mmol) in methanol (20 ml) was allowed to stand for 24 h at ambient temperature. The mixture was acidified with 1 M HCl solution (50 ml) and extracted with ethyl acetate (4 x 15 ml). The combined extracts were washed with saturated NaCl solution (1 x 10 ml) and dried (MgSO<sub>4</sub>). After filtration and evaporation, the residue was purified by flash chromatography on a 100 x 65 mm silica gel 60 column eluted with ethyl acetate-heptane (1:1), taking 50-ml fractions, giving 1.3 g (66%).

<sup>1</sup>H NMR (200 MHz; CDCl<sub>3</sub>): δ 0.88 (3H, t, J = 6.6 Hz), 1.25-1.5 (4H, m), 1.95-2.05 (2H, m), 6.49 (1H, br s), 6.75 (2H, d, J = 8.9 Hz), 7.12 (2H, d, J = 8.9 Hz), 7.1-7.35 (5H, m).

<sup>13</sup>C NMR (50 MHz; CDCl<sub>3</sub>): δ 13.6, 22.3, 24.3, 36.2, 72.8, 114.3, 121.9, 123.8, 125.3, 125.7, 127.2, 133.5, 154.6, 169.0, 169.5.

**EXAMPLE 8****Preparation of 4-(1-Butyl)-4-hydroxy-1-(4-**

- 17 -

methoxyphenyl)-2-phenyl-3,5-pyrazolidinedione

Prepared from 4-(1-butyl)-1-(4-methoxyphenyl)-2-phenyl-3,5-pyrazolidinedione (1.35 g; 3.8 mmol), 35% H<sub>2</sub>O<sub>2</sub> (4.3 ml; 50 mmol), 2 M NaOH (0.35 ml; 0.7 mmol), and methanol (50 ml) using the procedure of Example 7. Purified by flash chromatography on a 110 × 65 mm silica gel 60 column eluted with ethyl acetate-heptane (1:1) to give 0.7 g (52%).

<sup>1</sup>H NMR (200 MHz; CDCl<sub>3</sub>): δ 0.85 (3H, t, J = 6.2 Hz), 1.2-1.5 (4H, m), 2.0-2.1 (2H, m), 3.69 (3H, s), 4.8 (1H, br s), 6.77 (2H, d, J = 9.0 Hz), 7.19 (2H, d, J = 9.0 Hz), 7.1-7.35 (5H, m).

<sup>13</sup>C NMR (50 MHz; CDCl<sub>3</sub>): δ 13.5, 22.3, 24.3, 36.7, 54.7, 73.3, 113.0, 122.1, 123.8, 125.8, 126.1, 127.5, 133.0, 156.7, 168.5, 169.0.

EXAMPLE 9Preparation of 1,2-Diphenyl-4-hydroxy-4-[2-(phenylsulfonyl)ethyl]-3,5-pyrazolidinedione

Prepared from (±)-sulfinpyrazone (2.02 g; 5.0 mmol), 35% H<sub>2</sub>O<sub>2</sub> (4.3 ml; 50 mmol), 2 M NaOH (0.35 ml; 0.7 mmol), and methanol (50 ml) using the procedure of Example 7. Purified by flash chromatography on a 130 × 65 mm silica gel 60 column eluted with ethyl acetate-acetic acid (20:1) to give 80 mg (4%).

<sup>1</sup>H NMR (200 MHz; CDCl<sub>3</sub>): δ 2.1-2.5 (2H, m), 3.0-3.7 (2H, m), 5.5 (1H, br s), 6.4-7.9 (15H, m).

<sup>13</sup>C NMR (50 MHz; CDCl<sub>3</sub>): δ 28.7, 47.5, 70.2, 121.5, 122.9, 125.8, 126.5, 127.4, 127.7, 127.9, 129.9, 133.3, 133.4, 136.9, 139.2, 167.5, 168.0.

EXAMPLE 101,2-Diphenyl-4-hydroxy-4-(4-methylphenyl)-3,5-

**pyrazolidinedione**

A mixture of 1,2-diphenyl-4-(4-methylphenyl)-3,5-pyrazolidinedione (1.10 g; 3.2 mmol), 35% H<sub>2</sub>O<sub>2</sub> (0.47 ml; 5.5 mmol), and acetic acid (40 ml) was stirred 16 days at room temperature. Sodium metabisulfite (1.0 g) was added and excess acetic acid evaporated off. The residue was dissolved in hot ethyl acetate (25 ml) and benzene (25 ml) and filtered. After cooling to room temperature, the mixture was filtered and the residue recrystallized from 50% aqueous ethanol (20 ml) to give 0.58 g (53%).

<sup>1</sup>H NMR (200 MHz; CDCl<sub>3</sub>): δ 2.32 (3H, s), 7.0-7.45 (14H, m).

<sup>13</sup>C NMR (50 MHz; CDCl<sub>3</sub>): δ 21.1, 57.9, 123.9, 124.5, 127.0, 128.3, 128.4, 128.7, 130.4, 135.6, 139.0, 168.5.

**EXAMPLE 11****Preparation of 4-Benzyl-1,2-diphenyl-4-hydroxy-3,5-pyrazolidinedione**

Prepared from 4-benzyl-1,2-diphenyl-3,5-pyrazolidinedione (3.3 g; 9.6 mmol), 35% H<sub>2</sub>O<sub>2</sub> (1.4 ml; 16.3 mmol), and acetic acid (50 ml) using the procedure of Example 10 to give 1.0 g (30%).

<sup>1</sup>H NMR (200 MHz; CDCl<sub>3</sub>): δ 3.30 (2H, s), 6.75-7.3 (15H, m).

<sup>13</sup>C NMR (50 MHz; CDCl<sub>3</sub>): δ 43.1, 75.4, 123.0, 126.7, 127.5, 128.4,, 130.2, 132.1, 134.7, 170.1.

**EXAMPLE 12****Antiviral activity of 4OH-OPB (Example 7)**

4OH-OPB was added to cultures of growing MT4 cells (a

- 19 -

human CD4 cell line). HIV-1, stored in the culture medium at  $-75^{\circ}\text{C}$  was thawed and added in an amount which infected about 1 in 7 cells in each culture. The virus was absorbed to the cells for 2.3 hours at ambient temperature whereafter the cultures were centrifuged at 1200 rpm, the medium was removed, the cells were suspended in fresh growth medium and 4OH-OPB was added to concentrations of 1, 10 and 100  $\mu\text{M}$  (diluted in medium from a stock solution of 20 mM in DMSO). After 72 hours the HIV antigen concentration was determined using Abbott's test. By way of comparison phenbutazone (PB) was tested analogously. The results are shown in Figure 1 and demonstrate inhibition of virus production by 4OH-OPB at concentrations above the lowest tested.

### EXAMPLE 13

#### Combination Antiviral effect with 4OH-OPB

Cell culture experiments were carried out as in Example 12, but in place of 4OH-OPB (0-100  $\mu\text{M}$ ) was added:

- i) 4OH-OPB (0-10  $\mu\text{M}$ ) with AZT (0-1  $\mu\text{M}$ )
- ii) 4OH-OPB (0-100  $\mu\text{M}$ ) with Indinavir (0-100  $\mu\text{M}$ )
- iii) 4OH-OPB (0-100  $\mu\text{M}$ ) with Nevirapin (0-10  $\mu\text{M}$ )
- iv) 4OH-OPB (0-10  $\mu\text{M}$ ) with ddI (0-100  $\mu\text{M}$ )

The results are shown in Figures 2-5 respectively and demonstrate the enhanced anti-HIV effect of 4OH-OPB in combination with other anti-viral agents.

### EXAMPLE 14

#### Preparation of capsules for oral use

4-OH OPB (Example 7)	50 mg
Amylum maydis	q.s.

The powder is mixed and filled into hard gelatin capsules (Capsugel size 00).

#### EXAMPLE 15

##### Preparation of tablets

	Gram
4-OH OPB (Example 7)	200
Lactose	85
Polyvinylpyrrolidone	5
Starch	42
Talcum powder	15
Magnesium stearate	3

4-OH OPB and lactose are screened through a 0.15 mm sieve and mixed together with an aqueous solution of polyvinyl-pyrrolidone. The mass is granulated, and the dried (40°C) granulate is mixed with starch, talcum powder and magnesium stearate. The granulate is compressed into tablets. The tablet diameter is 11 mm, the tablet weight is 350 mg and each tablet contains 200 mg 4-OH OPB.

#### EXAMPLE 16

##### Preparation of a suspension for rectal administration

Methyl p-hydroxybenzoate (70 mg) and propyl-p-hydroxybenzoate (15 mg) are dissolved in water (100 ml) at 90°C. After cooling to 30°C, methyl cellulose (2g) is added and the mixture is agitated for 3 hours. 1 gram 4-OH OPB (Example 7) is screened through a 0.15 mm sieve, and dispersed in the solution under vigorous stirring. The suspension is filled in a 100 ml tube. The suspension contains 10 mg 4-OH OPB/ml.

- 21 -

EXAMPLE 17Preparation of oral suspension

	Gram
4OH OPB (Example 7)	10
Carboxymethyl cellulose	1.5
Sorbitol	200
Sodium benzoate	1.0
Orange essence	0.3
Apricot essence	0.7
Ethanol	50
Water	236.5

Carboxymethyl cellulose, sorbitol and sodium benzoate are dissolved in water with stirring for 2 hours. A solution of the essences in ethanol is added. 4-OH OPB is screened through a 0.15 mm sieve and dispersed in the solution under vigorous stirring. The suspension (10 gram) is filled in a 20 ml tube. Each tube contains 200 mg 4-OH OPB.

EXAMPLE 18Mouse toxicity

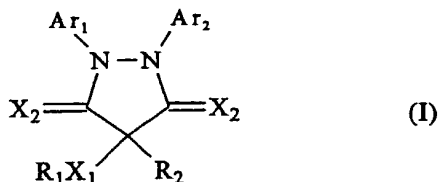
20g mice were given single doses of 4OH-OPB (20 mM in DMSO) intraperitoneally. Doses of 1 to 100  $\mu$ M (in ECF), corresponding to 0.29 to 29  $\mu$ M/kg bodyweight, produced no toxic effect. Furthermore, injection of 4OH-OPB could be increased to 2000 mg/kg (corresponding to 20 mM in the extracellular fluid) before the mice started to die (6 out of 10 died at 2000 mg/kg). Thus the concentrations that effectively inhibit HIV replication in cell cultures are up to 200000 times lower than the lethal dose in mice.



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Claims

1. The use of a compound of formula I



(where each  $X_2$ , which may be the same or different is O or S,

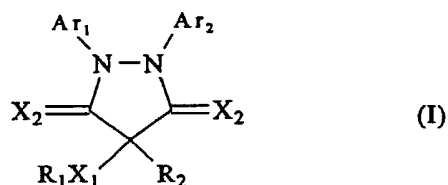
$X_1$  is O, OO or S,

$R_1$  is hydrogen or a hydroxyl or thiol protecting group,  
 $R_2$  is hydrogen or a alkyl, alkenyl, alkynyl, alkaryl, aralkyl or aralkenyl group, containing up to 10 carbons, optionally substituted by a sulphonyl group, and each of

$Ar_1$  and  $Ar_2$ , which may be the same or different, is a homo or heterocyclic aromatic group comprising 5 to 7 membered aromatic ring, optionally carrying a fused aromatic ring and optionally substituted on ring atoms by  $C_{1-6}$  alkyl, hydroxy, thiol,  $C_{1-6}$  alkoxy, cyano, Cl, F, Br, I, protected hydroxy, or protected thiol), or a physiologically acceptable salt thereof, for the manufacture of a medicament for use in therapy or prophylaxis.

2. A method of treatment of the human or non-human body to combat an inflammatory or viral disease, which method comprises administering to said body a compound of formula I

- 23 -



(where each  $X_2$ , which may be the same or different is O or S,

$X_1$  is O, OO or S,

$R_1$  is hydrogen or a hydroxyl or thiol protecting group,

$R_2$  is hydrogen or a alkyl, alkenyl, alkynyl, alkaryl, aralkyl or aralkenyl group, containing up to 10 carbons, optionally substituted by a sulphonyl group, and each of  $Ar_1$  and  $Ar_2$ , which may be the same or different, is a homo or heterocyclic aromatic group comprising 5 to 7 membered aromatic ring, optionally carrying a fused aromatic ring and optionally substituted on ring atoms by  $C_{1-6}$  alkyl, hydroxy, thiol,  $C_{1-6}$  alkoxy, cyano, Cl, F, Br, I, protected hydroxy, or protected thiol), or a physiologically acceptable salt thereof.

3. A method as claimed in claim 2 comprising administering said compound, or a physiologically acceptable salt thereof in combination with another antiviral agent.

4. A method as claimed in claim 3 wherein said additional antiviral agent is at least one antiviral agent selected from a reverse transcriptase inhibitor and a protease inhibitor.

5. A method as claimed in claim 3 wherein said additional antiviral agent is an agent selected from the group of AZT, indinavir, nevirapine and 2',3'-dideoxyinosine (ddI).

- 24 -

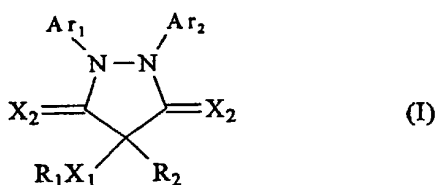
6. A method as claimed in any of claims 2 to 5 wherein said disease is a disease caused by a pathogen from the group of togaviridea, reoviridea, picornaviridea, hantaviridea, orthomyxoviridea, paramyxoviridea, mononegaviralis, viral hepatitis, haemorrhagic fevers, flaviviridea, viral encephalitis, coronaviridea, calciviridea, adenoviridea, papovaviridea, arboviridea, pox virus, rhabdoviridea, arenaviridea HIV-1, HIV-2, HTLV-I, HTLV-II and herpes viruses.

7. A method of combatting HIV infection which comprises administering to an HIV-infected patient a T-lymphocyte growth suppressing agent in an amount sufficient to suppress T-lymphocyte growth in said patient for a period sufficient to reduce the T-lymphocyte concentration in the lymphatic system in said patient by at least 25% said administration being repeated at intervals of at least 3 months.

8. A method of combatting HIV infection as claimed in claim 7 wherein said T-lymphocyte growth suppressing agent is a pyrazolidinol.

9. A method as claimed in claim 7 or claim 8 wherein said interval is at least 9 months.

10. A method as claimed in any of claims 7 to 9 wherein a compound of formula I



(where each  $\text{X}_2$ , which may be the same or different is O or S,

Art. 34

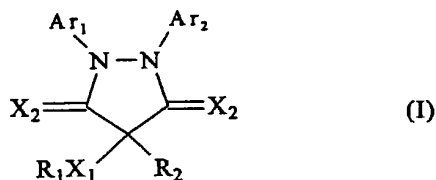
- 25 -

$X_1$  is O, OO or S,

$R_1$  is hydrogen or a hydroxyl or thiol protecting group,

$R_2$  is hydrogen or a alkyl, alkenyl, alkynyl, alkaryl, aralkyl or aralkenyl group, containing up to 10 carbons, optionally substituted by a sulphonyl group, and each of  $Ar_1$  and  $Ar_2$ , which may be the same or different, is a homo or heterocyclic aromatic group comprising 5 to 7 membered aromatic ring, optionally carrying a fused aromatic ring and optionally substituted on ring atoms by  $C_{1-6}$  alkyl, hydroxy, thiol,  $C_{1-6}$  alkoxy, cyano, Cl, F, Br, I, protected hydroxy, or protected thiol), or a physiologically acceptable salt thereof is administered in a daily dose of 0.1 to 10  $\mu\text{mol/kg}$  bodyweight.

11. A pharmaceutical composition comprising a compound of formula I



(where each  $X_2$ , which may be the same or different is O or S,

$X_1$  is O, OO or S,

$R_1$  is hydrogen or a hydroxyl or thiol protecting group,

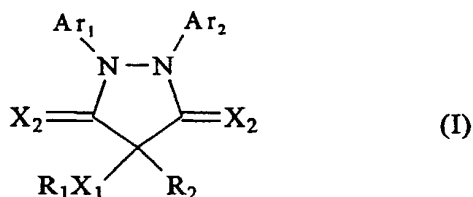
$R_2$  is hydrogen or a alkyl, alkenyl, alkynyl, alkaryl, aralkyl or aralkenyl group, containing up to 10 carbons, optionally substituted by a sulphonyl group, and each of  $Ar_1$  and  $Ar_2$ , which may be the same or different, is a homo or heterocyclic aromatic group comprising 5 to 7 membered aromatic ring, optionally carrying a fused aromatic ring and optionally substituted on ring atoms by  $C_{1-6}$  alkyl, hydroxy, thiol,  $C_{1-6}$  alkoxy, cyano, Cl, F, Br, I, protected hydroxy, or protected thiol), or a physiologically acceptable salt thereof, together with

- 26 -

at least one pharmaceutically acceptable carrier or excipient.

12. A pharmaceutical composition as claimed in claim 11 additionally comprising another antiviral agent.

13. A compound of formula I



(where each  $X_2$ , which may be the same or different is O or S,

$X_1$  is O, OO or S,

$R_1$  is hydrogen or a hydroxyl or thiol protecting group,

$R_2$  is hydrogen or a alkyl, alkenyl, alkynyl, alkaryl, aralkyl or aralkenyl group, optionally substituted by a sulphonyl group, and

one of  $Ar_1$  and  $Ar_2$  is Ph and the other is 4-hydroxyphenyl,

or a salt thereof, providing that if  $R_2$  is  $C_4H_9$ ,  $R_1$  is not H or OH.

14. A compound as claimed in claim 13 or claim 14 wherein one  $X_2$  group is S.

15. A compound as claimed in either of claims 13 or 14 wherein  $X_1$  is O.

16. A compound as claimed in any of claims 13 to 15 wherein  $R_1$  is acyl.

17. A compound as claimed in any of claims 13 to 16 wherein  $R_1$  is hydrogen.

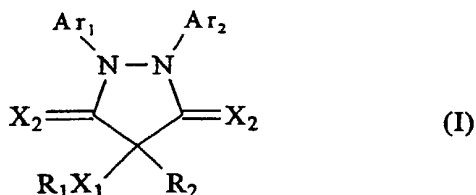
- 27 -

18. A compound as claimed in claim 13 wherein each  $X_2$  is oxygen,  $R_1X_1$  is HO or  $CH_3CO.O$ , and  $R_2$  is  $C_{1-6}$  alkyl or alkenyl, or a salt thereof.

19. A compound as claimed in any of claims 13 to 18 for use as a medicament.

20. 4-Butyl-4-hydroxy-2(p-hydroxyphenyl)-1-phenyl-3,5-pyrazolidinedione for use as a medicament.

21. A method of treatment of the human or non-human body to combat an autoimmune disease or tissue rejection, which method comprises administering to said body a compound of formula I



(where each  $X_2$ , which may be the same or different is O or S,

$X_1$  is O, OO or S,

$R_1$  is hydrogen or a hydroxyl or thiol protecting group,

$R_2$  is hydrogen or a alkyl, alkenyl, alkynyl, alkaryl, aralkyl or aralkenyl group, containing up to 10 carbons, optionally substituted by a sulphonyl group, and each of  $Ar_1$  and  $Ar_2$ , which may be the same or different, is a homo or heterocyclic aromatic group comprising 5 to 7 membered aromatic ring, optionally carrying a fused aromatic ring and optionally substituted on ring atoms by  $C_{1-6}$  alkyl, hydroxy, thiol,  $C_{1-6}$  alkoxy, cyano, Cl, F, Br, I, protected hydroxy, or protected thiol) or a physiologically tolerable salt thereof.

- 28 -

22

21. A method of claim 21 wherein said disease is selected from Addison's disease, Behçet's syndrome, diabetes mellitus, haemolytic anaemia, lupus erythematosus, multiple sclerosis, myasthenia gravis, pernicious anaemia, polyglandular deficiency, polymyositis, dermatomyositis, testicular failure, thrombocytopenic purpura, Crohns disease, ulcerative colitis and rheumatoid arthritis.

23

24. A method of claim 21 wherein said tissue rejection is tissue rejection following transplant.

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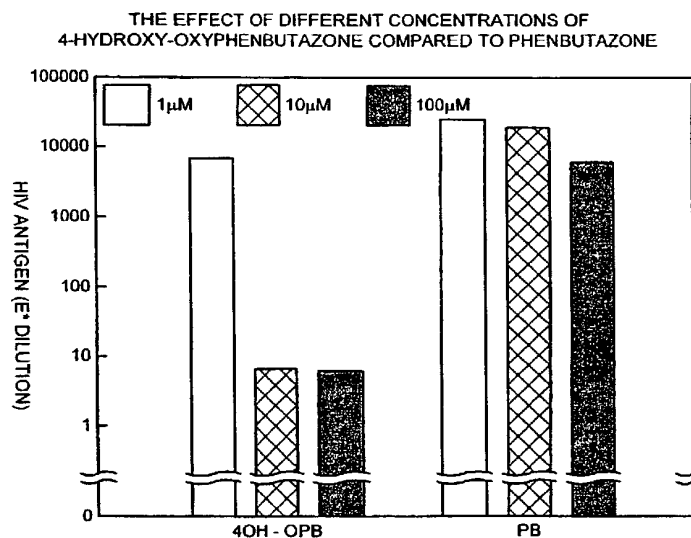
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[Continued on next page]

(54) Title: **PYRAZOLIDINOL COMPOUNDS**



(57) Abstract: The invention provides the use of an optionally hydroxy-protected 4-hydroxy or hydroperoxy-3,5-dioxypyrazolidine or an equivalent wherein a pyrazolidine ring attached oxygen is replaced by a sulphur, or a physiologically acceptable salt thereof, for the manufacture of a medicament for use in therapy or prophylaxis. Additionally, the invention provides a method of combating HIV infection which comprises administering to an HIV-infected patient a T-lymphocyte growth suppressing agent, preferably a pyrazolidinol, in an amount sufficient to suppress T-lymphocyte growth in said patient for a period sufficient to reduce the T-lymphocyte concentration in lymph nodes in said patient by at least 25 % said administration being repeated at intervals of at least 3 months.

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**Published:**

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- Before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments.

*For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.*

FIG. 1

THE EFFECT OF DIFFERENT CONCENTRATIONS OF  
4-HYDROXY-OXYPHENBUTAZONE COMPARED TO PHENBUTAZONE

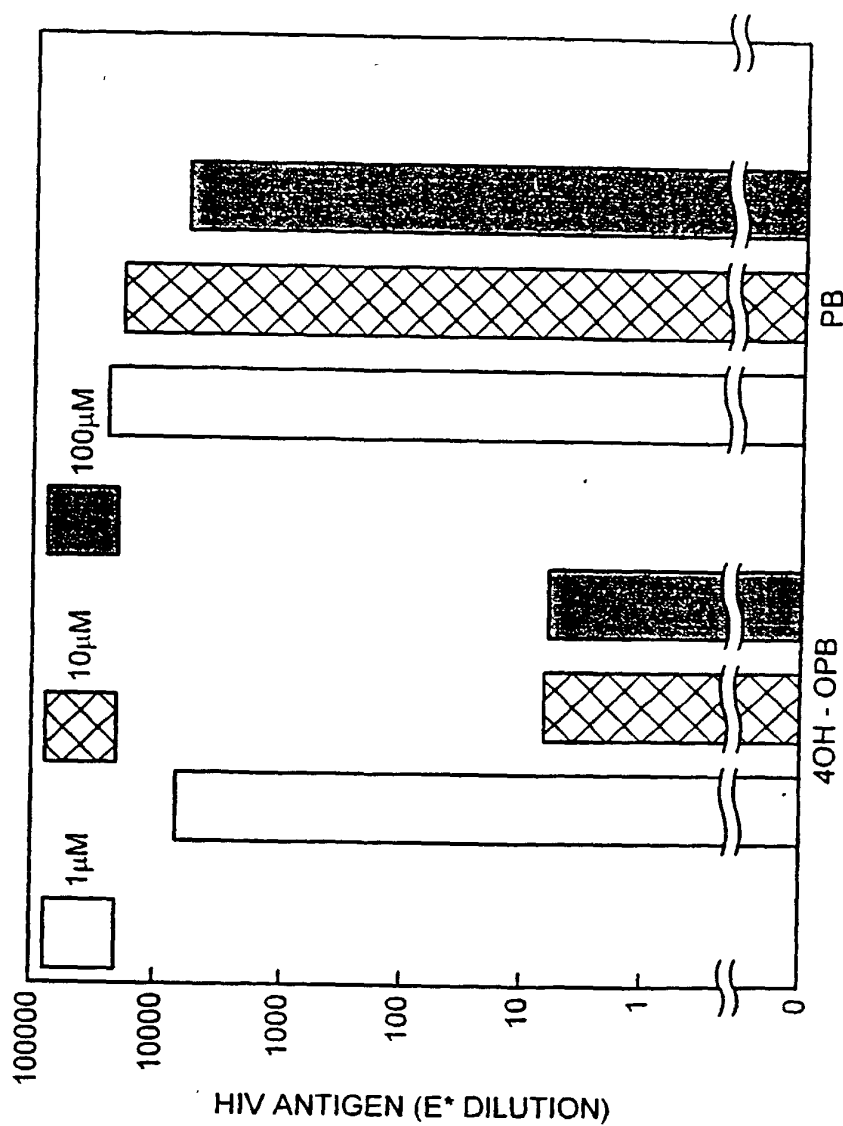
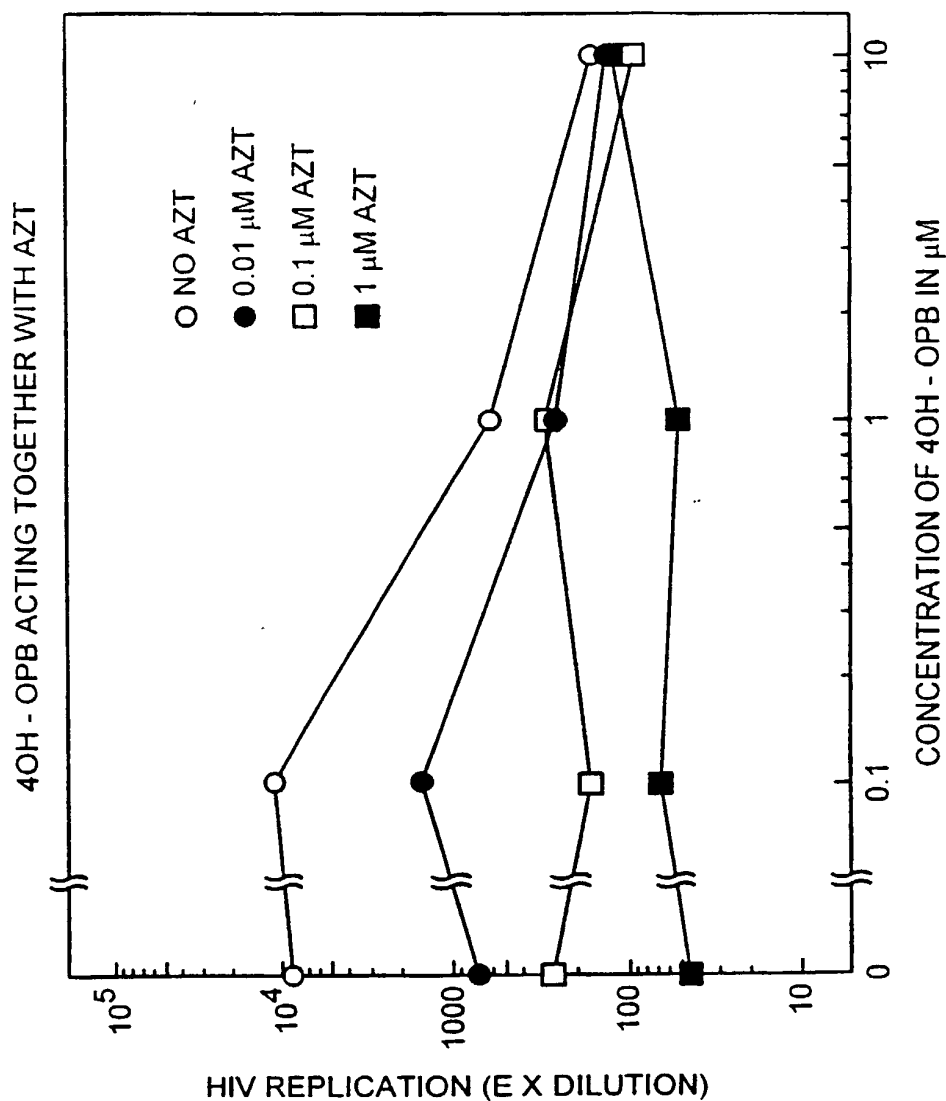


FIG. 2



3 / 5

FIG. 3

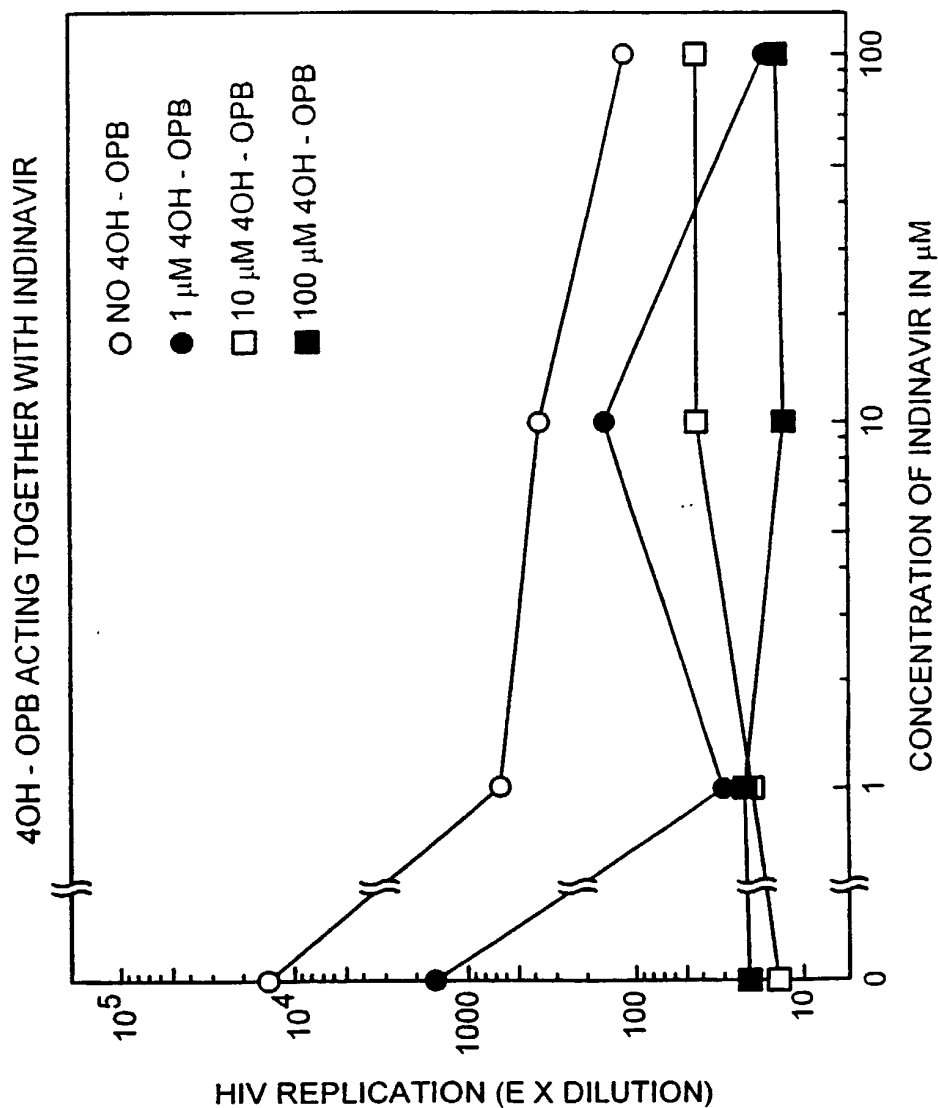
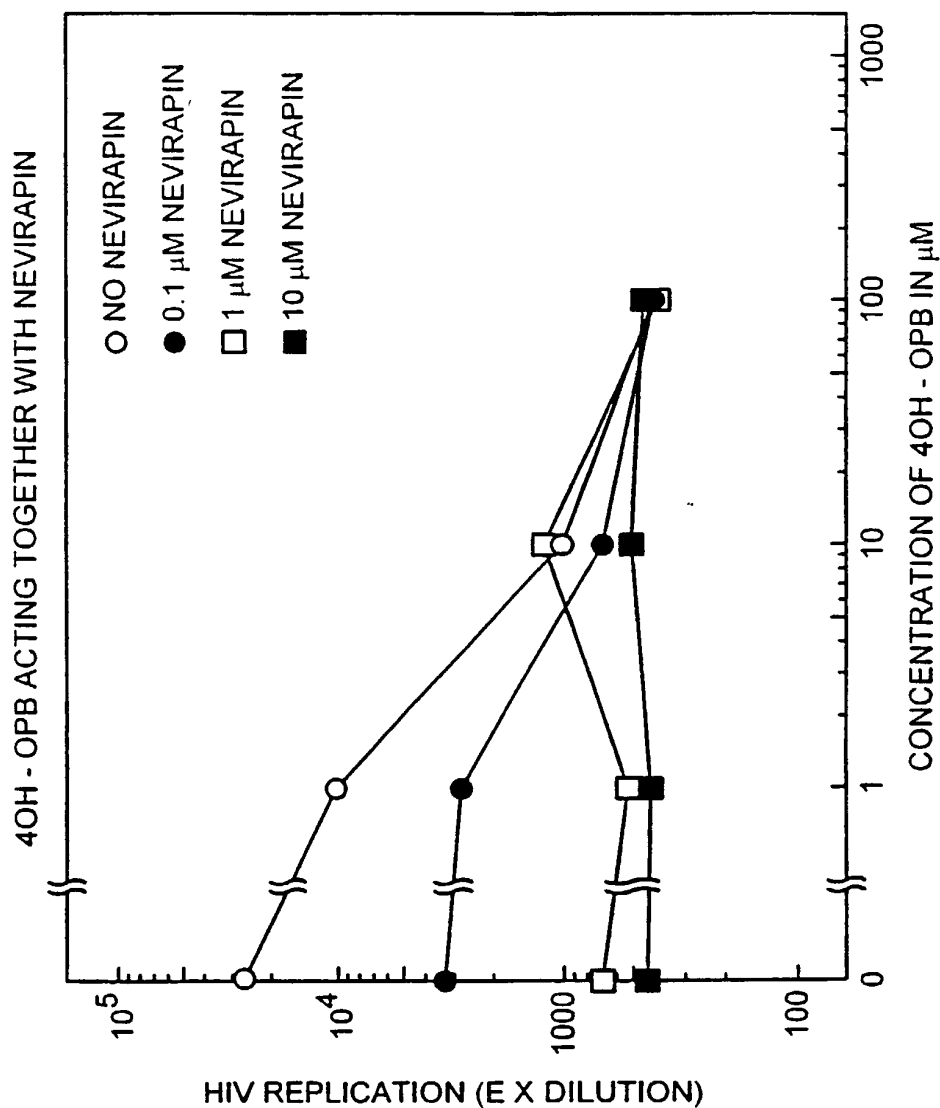
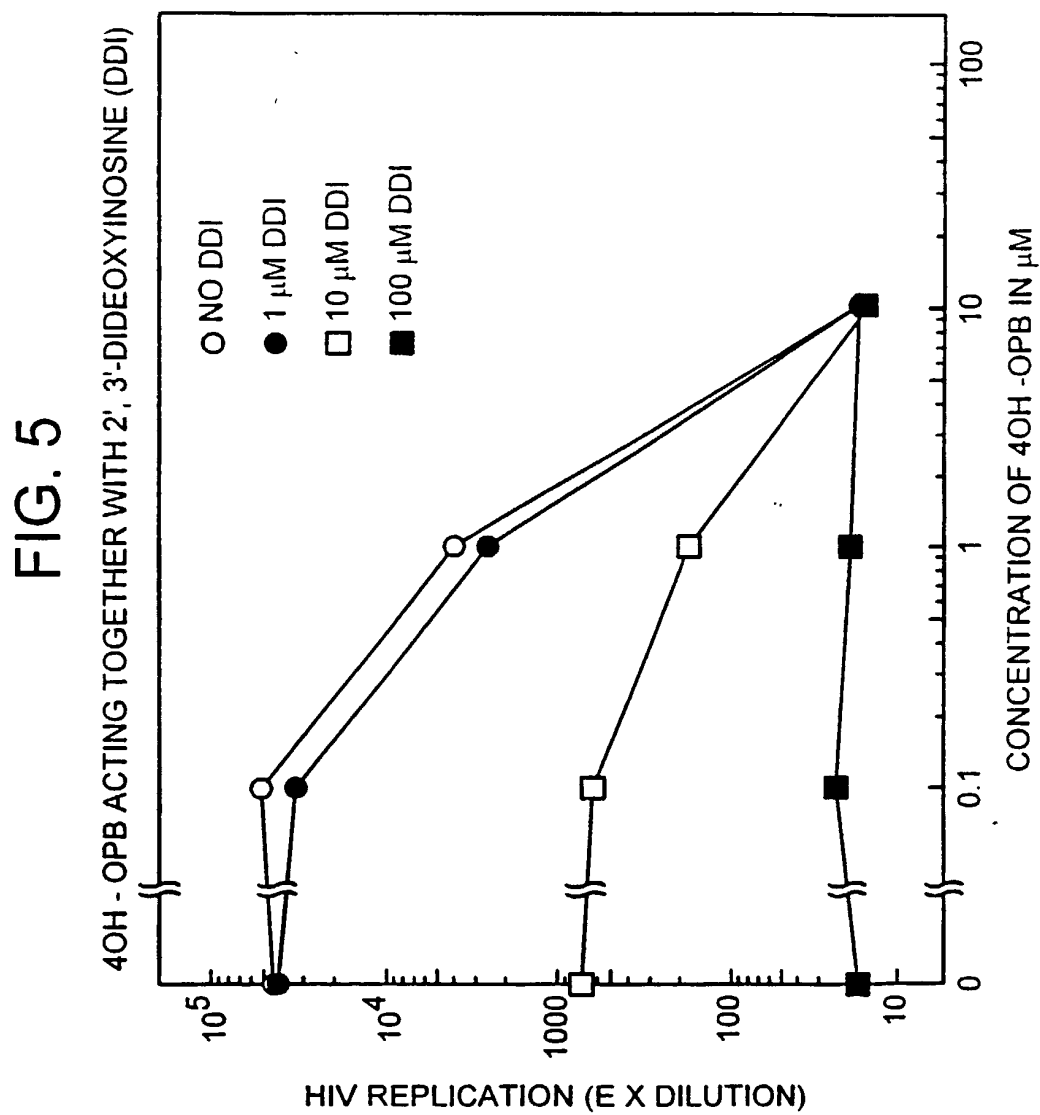


FIG. 4





**DECLARATION AND POWER OF ATTORNEY**  
**For Utility or Design Application**

As a below named inventor, I hereby declare that my residence, post office address and citizenship are as stated below next to my name.

I believe I am the original, first, and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed for and for which a patent is sought on the invention entitled:

**PYRAZOLIDINOL COMPOUNDS**

the specification of which is attached hereto **OR** was filed on June 29, 2000 as United States Application Number \_\_\_\_\_ or PCT International Application Number PCT/GB00/02513 and was amended on December 28, 2001 (if applicable).

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment specifically referred to above.

I acknowledge the duty to disclose information which is material to patentability as defined in 37 CFR § 1.56, including for continuation-in-part applications, material information which became available between the filing date of the prior application and the national or PCT international filing date of the continuation-in-part application.

I hereby claim foreign priority benefits under 35 U.S.C. § 119(a)-(d) or (f), or 365(b) of any foreign application(s) for patent, inventor's or plant breeder's rights certificate(s), or 365(a) of any PCT international application which designated at least one country other than the United States of America, listed below and have also identified below, by checking the box, any foreign application for patent, inventor's or plant breeder's rights certificate(s), or any PCT international application having a filing date before that of the application on which priority is claimed.:

Country	Application Number	Date of Filing	Priority Claimed Under 35 U.S.C. § 119	
			YES	NO
Great Britain	9915184.7	June 29, 1999	X	

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Residence	Citizenship	
Post Office Address		